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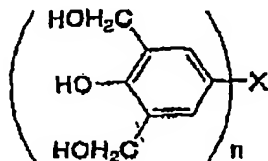
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(54) **Hydroxymethyl-substituted polyfunctional phenols**

(57) Hydroxymethyl-substituted polyfunctional phenols, expressed by the following structure:

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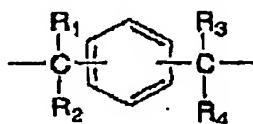
General structure (I)



(I)

(wherein X represents: bivalent group (a) expressed by the following structure:

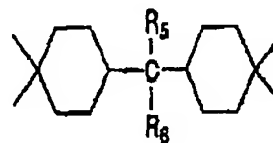
General structure (II)



(II)

(wherein R₁, R₂, R₃ and R₄ each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);
quadrivalent group (b) expressed by the following struc-

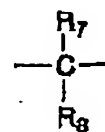
General structure (III)



(III)

(wherein R₅ and R₆ each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);
or bivalent group (c) expressed by the following structure:

General structure (IV)



(IV)

(wherein R₇ and R₈ each independently represent a hy-

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drogen atom or a monofluoromethyl, difluoromethyl or trifluoromethyl group; however, R_7 and R_8 cannot be both hydrogen atoms); wherein n takes 2 when X is bi-

valent group (a), takes 4 when X is quadrivalent group (b), or takes 2 when X is bivalent group (c)).

Description

[0001] This invention relates to new hydroxymethyl-substituted polyfunctional phenols, wherein one phenol nucleus contains two hydroxymethyl groups as nucleus-substitution groups. Such compounds may be used as a modifier for phenol resins, etc., photo-resist material, or material for producing various polyphenol compounds involving chemical reaction of said compounds with phenols.

[0002] The publication of Japanese Patent Application Laid-open No. 53-71044 discloses bis (4-hydroxy-3,5-dihydroxymethyl phenyl) methane, which is a bisphenol compound containing hydroxymethyl groups as nucleus-substitution groups. The publication of Japanese Patent Application Laid-open No. 8-277235 also discloses 3,3',5,5'-tetrahydroxymethyl-4,4'-diphenol as an intermediate material used for producing polyphenols via chemical reaction with phenols.

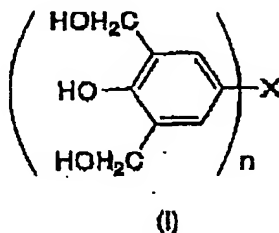
[0003] Among bisphenol compounds containing hydroxymethyl groups as nucleus-substitution groups, such as those mentioned above, polyfunctional phenols having an aromatic ring (e.g. benzene ring) or cycloalkane ring (e.g. cyclohexane ring) at the center of symmetry provide a high melting point. Polyphenol compounds made from these polyfunctional phenols are also expected to exhibit excellent resistance to heat. In addition, polyfunctional phenols having a fluorine-substituted alkylene group at the center of symmetry of the molecule are expected to offer property-modifying effects, such as improving the water repellency of phenol resins.

[0004] However, hydroxymethyl-substituted polyfunctional phenols, wherein the molecule of said phenol contains two or four phenol nuclei each having two hydroxymethyl groups as nucleus-substitution groups and a benzene ring, cyclohexane ring or fluorine-substituted alkylene group at the center of symmetry of the molecule, have not heretofore been known.

[0005] In light of the aforementioned condition to date regarding hydroxymethyl-substituted polyfunctional phenols containing hydroxymethyl groups as nucleus-substitution groups, the present invention aims to provide new hydroxymethyl-substituted polyfunctional phenols, wherein the molecule of said phenol contains two or four phenol nuclei each having two hydroxymethyl groups as nucleus-substitution groups and a benzene ring, cyclohexane ring or fluorine-substituted alkylene group at the center of symmetry of the molecule.

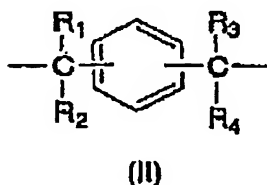
[0006] The present invention provides hydroxymethyl-substituted polyfunctional phenols expressed by the following structure:

General structure (I)



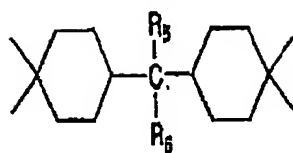
(wherein X represents: bivalent group (a) expressed by the following structure:

General structure (II)



(wherein R₁, R₂, R₃ and R₄ each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4); quadrivalent group (b) expressed by the following structure;

General structure (III)



(III)

(wherein R₅ and R₆ each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);

or bivalent group (c) expressed by the following structure:



(IV)

(wherein R₇ and R₈ each independently represent a hydrogen atom or a monofluoromethyl, difluoromethyl or trifluoromethyl group; however, R₇ and R₈ cannot be both hydrogen atoms);
wherein n takes 2 when X is bivalent group (a), takes 4 when X is quadrivalent group (b), or takes 2 when X is bivalent group (c).

[0007] The first type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention reflects aforementioned general structure (I), wherein X is bivalent group (a) expressed by aforementioned general structure (II). If R₁, R₂, R₃ and R₄ in bivalent group (a) are each an alkyl group with a carbon atom number of 1 through 4, specific examples of such alkyl groups include methyl, ethyl, propyl and butyl groups. Among these alkyl groups, propyl and butyl groups may have a linear or branched chain.

[0008] Therefore, 1,4-phenylenebis (methylidene) group, 1,4-phenylenebis (1-ethylidene) group, 1,4-phenylenebis (1-propylidene) group, 1,4-phenylenebis (1-butylidene) group, 1,4-phenylenebis (2-methyl-1-propylidene) group, 1,4-phenylenebis (1-pentylidene) group, 1,4-phenylenebis (3-methyl-1-butylidene) group, 1,4-phenylenebis (3-methyl-1-butylidene) group, 1,4-phenylenebis (2,2-dimethyl-1-propylidene) group, 1,4-phenylenebis (1-methyl ethylidene) group, 1,4-phenylenebis (1-methyl propylidene) group, 1,4-phenylenebis (1-methyl butylidene) group, 1,4-phenylenebis (1-methyl pentylidene) group, 1,4-phenylenebis (1-ethyl propylidene) group, 1,4-phenylenebis (1-ethyl butylidene) group, 1,4-phenylenebis (1-ethyl pentylidene) group, 1,4-phenylenebis (1-propyl butylidene) group, 1,4-phenylenebis (1-isopropyl-2-methyl propylidene) group, 1,4-phenylenebis (1-propyl pentylidene) group, 1,4-phenylenebis (1-butyl pentylidene) group, 1,4-phenylenebis (1-t-butyl-2,2-dimethyl propylidene) group, 1,3-phenylenebis (methylidene) group, 1,3-phenylenebis (1-methyl ethylidene) group, 1,3-phenylenebis (1-ethyl propylidene) group, 1,3-phenylenebis (1-propyl butylidene) group and 1,3-phenylenebis (1-butyl pentylidene) group are given as specific examples of bivalent group (a) expressed by aforementioned general structure (II).

[0009] Therefore, 4,4'-[1,4-phenylenebis (methylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-ethylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-propylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-pentylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-ethyl propylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-propyl butylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,4-phenylenebis (1-butyl pentylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,3-phenylenebis (methylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,3-phenylenebis (1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,3-phenylenebis (1-ethyl propylidene)] bis (3,5-dihydroxymethyl phenol), 4,4'-[1,3-phenylenebis (1-propyl butylidene)] bis (3,5-dihydroxymethyl phenol) and 4,4'-[1,3-phenylenebis (1-butyl pentylidene)] bis (3,5-dihydroxymethyl phenol) are given as specific examples of the first type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention.

[0010] The second type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention reflects aforementioned general structure (I), wherein X is quadrivalent group (b) expressed by aforementioned general structure (III). If R_5 and R_6 in quadrivalent group (b) are each an alkyl group with a carbon atom number of 1 through 4, specific examples of such alkyl groups include methyl, ethyl, propyl and butyl groups. Among these alkyl groups, propyl and butyl groups may have a linear or branched chain.

[0011] Therefore, methylenedioxy group, 1-ethylenedioxy group, 1-propylenedioxy group, 1-butylenedioxy group, 2-methyl-1-propylenedioxy group, 1-pentylenedioxy group, 3-methyl-1-butylenedioxy group, 2,2-dimethyl-1-propylenedioxy group, 1-methyl ethylenedioxy group, 1-methyl propylenedioxy group, 1-methyl butylenedioxy group, 1-methyl pentylenedioxy group, 1-ethyl propylenedioxy group, 1-ethyl butylenedioxy group, 1-ethyl pentylenedioxy group, 1-propyl butylenedioxy group, 1-isopropyl-2-methyl propylenedioxy group, 1-propyl pentylenedioxy group, 1-butyl pentylenedioxy group and 1-*t*-butyl-2, 2-dimethyl propylenedioxy group are given as specific examples of quadrivalent group (b) expressed by aforementioned general structure (III).

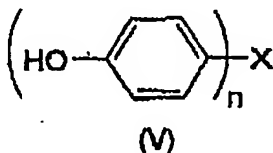
[0012] Therefore, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)] methane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)] ethane, 2,2-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)] propane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-methyl propane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-ethyl propane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-propyl butane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-isopropyl-2-methyl propane, 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-butyl pentane and 1,1-bis [4,4-cyclohexylenedioxybis (4-hydroxy-3,5-dihydroxymethyl phenyl)]-1-*t*-butyl-2,2-dimethyl propane are given as specific examples of the second type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention.

[0013] Further, the third type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention reflects aforementioned general structure (I), wherein X is bivalent group (c) expressed by aforementioned general structure (IV). R_7 and R_8 in bivalent group (c) are each an independent hydrogen atom or a monofluoromethyl group, difluoromethyl group or trifluoromethyl group. However, R_7 and R_8 cannot be both hydrogen atoms.

[0014] Therefore, 2-fluoroethylenedioxy group, 2-difluoroethylenedioxy group, 2-trifluoroethylenedioxy group, 1-fluoromethyl-2-fluoroethylenedioxy group, 1-fluoromethyl-2-difluoroethylenedioxy group, 1-difluoromethyl-2-difluoroethylenedioxy group and 1-trifluoromethyl-2-trifluoroethylenedioxy group are given as specific examples of aforementioned bivalent group (c).

[0015] Therefore, 4,4'-(2-fluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol), 4,4'-(2-difluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol), 4,4'-(2-trifluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol), 4,4'-(1-fluoromethyl-2-fluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol), 4,4'-(1-difluoromethyl-2-difluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol) and 4,4'-(1-trifluoromethyl-2-trifluoroethylenedioxy) bis (3,5-dihydroxymethyl phenol) are given as specific examples of the third type of new hydroxymethyl-substituted polyfunctional phenols provided by the present invention.

[0016] The hydroxymethyl-substituted polyfunctional phenols provided by the present invention, which are expressed by aforementioned general structure (I), may be obtained from multivalent phenol compounds expressed by general structure (V) below:



(wherein X and n are the same as defined earlier).

To be specific, each multivalent phenol compound of the above structure is reacted with 1 to 10 mols, or preferably 1.5 to 5 mols, of formaldehyde per 1 mol of hydroxymethyl group contained in 1 mol of the target substance, in the presence of basic catalyst or in water or a mixed solvent consisting of water and organic solvent, after which the obtained reaction product is neutralized,

[0017] The amount of formaldehyde will significantly exceed the theoretical value in this reaction, if more than 10 mols of formaldehyde is reacted with the aforementioned multivalent phenol compound per 1 mol of hydroxymethyl group contained in 1 mol of the target substance, and therefore the process economy will drop. On the other hand, it is equally undesirable to react less than 10 mols of formaldehyde with the aforementioned multivalent phenol compound per 1 mol of hydroxymethyl group contained in 1 mol of the target substance, since a significant amount of monohydroxymethyl-group substitution products or unreacted bisphenols will remain.

[0018] The formaldehyde used in the aforementioned production method may be a commercially available, undiluted formalin solution. Alternatively, paraformaldehyde or trioxane that reacts in a similar manner to formaldehyde in water can also be used. However, use of formalin is preferred.

[0019] The aforementioned basic catalyst may be a hydroxide of alkali metal such as sodium hydroxide, potassium hydroxide or lithium hydroxide, acetate, calcium hydroxide, or a bivalent metal such as zinc hydroxide or magnesium hydroxide. However, use of a hydroxide of alkaline earth, acetate, pyridine, or a tertiary amine such as trimethyl amine or tributyl amine is preferred. Among these, sodium hydroxide or potassium hydroxide is most preferred,

5 [0020] In the present invention, such basic catalyst may be added by an equivalent of 0.1 to 5 times, or preferably 0.1 to 2 times, the hydroxyl amount in the aforementioned multivalent phenol compound. If the amount of basic catalyst exceeds 5 times the hydroxyl amount in the aforementioned multivalent phenol compound, the basic catalyst becomes excessive in the reaction system. This is not desirable, since an excessively large amount of acid will be required to acidify the reaction system for precipitation and collection of the reaction production.

10 [0021] In the present invention, the reaction of the above multivalent phenol compound and formaldehyde in the presence of basic catalyst is normally initiated in a water medium or a mixed solvent consisting of water and organic solvent. The amount of water/mixed solvent is normally around 1 to 5 times, or preferably 2 to 3 times, the weight of the reacting material, which, in this case, is the above multivalent phenol compound.

15 [0022] The aforementioned organic solvent may be an alcohol solvent such as methanol, ethanol, n-propyl alcohol, isopropyl alcohol, n-butanol, ethylene glycol, ethylene glycol monomethyl ether, diethylene glycol or carbitol, or a water-soluble organic solvent such as tetrahydrofuran, dioxane, dimethyl formamide, dimethyl acetamide or n-methyl pyrrolidone, as long as the solubility, in the mixed solvent, of the aforementioned basic catalyst and the reacting material, which, in this case, is the aforementioned multivalent phenol compound, will not be affected.

20 [0023] The above reaction is normally initiated in a temperature range of 0 to 60°C. or preferably 30 to 50°C, over a period of around 1 to 72 hours, or preferably 4 to 16 hours. A reaction temperature exceeding 60°C will produce large amounts of various unwanted byproducts, such as the substances with high molecular weight mentioned earlier.

25 [0024] The product obtained from said condensation reaction of the aforementioned multivalent phenol compound and formaldehyde in the presence of basic catalyst is a mixture containing said multivalent phenol compound having hydroxymethyl groups added to the aromatic nucleuses on both sides of its molecule. This molecular structure exists in the obtained reaction mixture partially or entirely as alkali salt, depending on the amount of basic catalyst used. This reaction is generally a solution reaction, although the specific form of reaction varies depending on the type and content of alcohol or other organic solvent in the water medium or mixed solvent used for reaction, amount of water medium/mixed solvent used, and type and amount of basic catalyst used.

30 [0025] Therefore, to separate and collect the target reaction product from the obtained reaction mixture, in the present invention the obtained reaction mixture is cooled and mixed with an extractant, which may be an organic, water-insoluble solvent such as any of the aromatic hydrocarbons, fatty alcohols, aliphatic ketones or esters, or any combination of such solvents. The reaction mixture with extractant is then added to an acid compound, such as organic acid, inorganic acid or water solution of such acid, to neutralize the entire reaction mixture and thereby separate the water layer and extract the reaction product into the organic layer. Next, thus obtained organic layer is condensed as necessary and then cooled to cause precipitation. Finally, the solid precipitate is filtered to obtain a hydroxymethyl-substituted polyfunctional phenol, which is the target substance.

35 [0026] In the present invention, an extractant can also be added after the reaction mixture is neutralized. In particular, the present invention provides a hydroxymethyl-substituted polyfunctional phenol of even higher purity by removing in advance the impurities separating from the reaction mixture using the aforementioned extractant in an early stage of neutralization, extracting the reaction product from the water layer into the organic layer using the aforementioned extractant, and then processing the organic layer as described above.

40 [0027] The aforementioned extractant may be any of the aromatic hydrocarbons such as benzene, toluene or xylene, fatty alcohols such as n-butanol, aliphatic ketones such as methyl isobutyl ketone or esters such as ethyl acetate or butyl acetate, which may be used alone or in combination.

45 [0028] Of the acid compounds that may be used in the aforementioned neutralization process, those that are organic acids include formic acid, acetic acid, propionic acid and oxalic acid. Inorganic acids include sulfuric acid, phosphoric acid, phosphorous acid, hypophosphorous acid and hydrochloric acid. Of these, use of hydrochloric acid and sulfuric acid is most preferred from the viewpoint of economy and ease of handling.

50 [0029] The chemical compounds obtained by the present invention using the method described above are normally in a solid state at normal temperature, and therefore can be used for various applications. In particular, the compounds provided by the present invention can be used as photo-resist material or to derive polyphenol compounds through further reaction with phenolic compounds. The compounds provided by the present invention are also usable as compounding agent that adds to the molecular weight of novolac phenol resins or as hardening agent for epoxy resins used in casting and powder coating. In these applications, the desired effect manifests after reaction with hydroxymethyl groups.

Examples

[0030] The present invention is explained by using examples. However, application of the present invention is not limited to the examples provided.

Example 1

Production of 4,4'-[1,4-phenylenebis (1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol)

[0031] 51.9 g (0.15 mol) of 4,4'-[1,4-phenylenebis (1-methyl ethylidene)] bisphenol, 10.8 g (0.45 mol) of lithium hydroxide and 176 g of pure water were mixed into a four-neck flask and agitated at room temperature for 10 minutes. Into the obtained solution kept at 40°C, 154 g (1.80 mol) of 35% formaldehyde solution was dropped to cause reaction over 2 hours. The reacted solution was kept at the same temperature for another 20 hours for thorough reaction,

[0032] At the end of the reaction period, the obtained reaction mixture (solution) was neutralized by adding 16% sulfuric acid, after which n-butyl alcohol was added to dissolve the reaction product. Thereafter, the water layer was separated and the obtained oil layer was rinsed in water. The oil layer was then distilled to obtain 77.2 g of the target substance in coarse state (63.5% purity as measured by high-speed liquid chromatography) as a light yellowish-brown solution. The solution was then refined by column chromatography to obtain the target 4,4'-[1,4-phenylenebis (1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol) as a white solid with 98% purity.

Mass spectrometry (LC-MS/AP): 466

Proton nuclear magnetic resonance spectrum (400 MHz, solvent: DMSO):

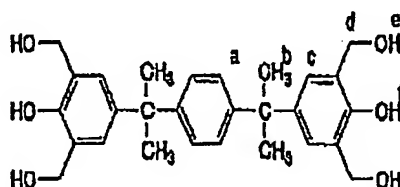


Table 1

δ (ppm)	Signal	Number of protons	Assignment
1.58	s	12	b
4.52	s	8	d
5.24	s	4	e
7.05	s	4	c
7.08	s	4	a
8.40	s	2	f

Example 2

Production of 4,4'-[1,3-phenylenebis(1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol)

[0033] 69.2 g (0.20 mol) of 4,4'-[1,3-phenylenebis (1-methyl ethylidene)] bisphenol, 150 g (0.60 mol) of 16% sodium hydroxide solution and 34.6 g of methanol were mixed into a four-neck flask and agitated at room temperature for 10 minutes. Into the obtained solution kept at 30°C, 205.7 g (2.4 mol) of 35% formaldehyde solution was dropped to cause reaction over 2 hours. The reacted solution was kept at the same temperature for another 7 hours for thorough reaction.

[0034] At the end of the reaction period, the obtained reaction mixture (solution) was neutralized by adding 16% sulfuric acid, after which n-butyl alcohol was added to dissolve the reaction product. Thereafter, the water layer was separated and the obtained oil layer was rinsed in water. The oil layer was then distilled to obtain 131 g of the target substance in coarse state (83.6% purity as measured by high-speed liquid chromatography) as a light yellow solution. The solution was then refined by column chromatography to obtain the target 4,4'-[1,3-phenylene bis (1-methyl ethylidene)] bis (3,5-dihydroxymethyl phenol) as a light brown solution with 98% purity.

Mass spectrometry (LC-MS/AP): 466

Proton nuclear magnetic resonance spectrum (400 MHz, solvent: DMSO):

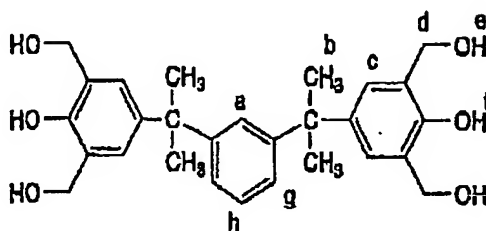


Table 2

δ (ppm)	Signal	Number of protons	Assignment
1.56	s	12	b
4.52	s	8	d
5.26	s	4	e
6.94	d	2	g
7.01	s	4	c
7.11	t	1	h
7.22	s	1	a
8.40	s	2	f

Example 3

Production of 2,2-bis [4,4-cyclohexylidenebis (4-hydroxy-3,5-dihydroxymethyl phenyl)] propane

[0035] 86.4 g (0.15 mol) of 2,2-bis [4,4-cyclohexylidenebis (4-hydroxy phenyl)] propane and 225 g (0.9 mol) of 16% sodium hydroxide solution were mixed into a four-neck flask and agitated at room temperature for 10 minutes. Into the obtained solution kept at 20°C, 321.4 g (3.75 mol) of 35% formaldehyde solution was dropped to cause reaction over approximately 1.7 hours. The reacted solution was kept at the same temperature for another 16.5 hours for thorough reaction.

[0036] At the end of the reaction period, the obtained reaction mixture (solution) was neutralized by adding 16% sulfuric acid, after which methyl ethyl ketone was added to dissolve the reaction product. Thereafter, the water layer was separated and the obtained oil layer was rinsed in water. The oil layer was then distilled to obtain 90 g of the target substance in coarse state (57, 1% purity as measured by high-speed liquid chromatography) as a white solid. The solid was dissolved in methanol and refined by column chromatography to obtain the target 2,2-bis [4,4-cyclohexylidenebis (4-hydroxy-3,5-dihydroxymethyl phenyl)] propane as a white solid with 93% purity.

Mass spectrometry (LC-MS/AP): 816

Proton nuclear magnetic resonance spectrum (400 MHz, solvent: DMSO):

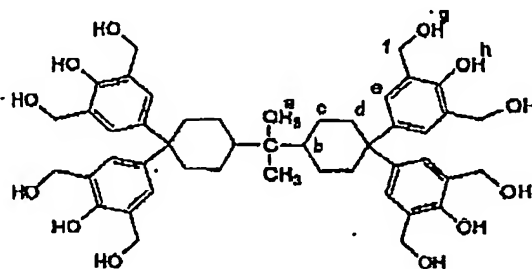


Table 3.

δ (ppm)	Signal	Number of protons	Assignment
0.47	s	6	a
1.09	t	4	c
1.33	t	2	b
1.53	d	4	c
1.69	t	4	d
2.69	d	4	d
4.46 to 4.53	d	16	f
5.19 to 5.27	t	8	g
6.99			
7.13	s	8	e
8.30			
8.34	s	4	h

Example 4

Production of 4,4'-(1-trifluoromethyl-2-trifluoroethylidene) bis (3,5-dihydroxymethyl phenol)

[0037] 168 g (0.5 mol) of 4,4'-(1-trifluoromethyl-2-trifluoroethylidene) bisphenol and 280.3 g (1.0 mol) of 20% potassium hydroxide solution were mixed into a four-neck flask and agitated at room temperature for 10 minutes. Into the obtained solution kept at 50°C, 342.9 g (4.0 mol) of 35% formaldehyde solution was dropped to cause reaction over approximately 1.5 hours. The reacted solution was kept at the same temperature for another 14,5 hours for thorough reaction.

[0038] At the end of the reaction period, the obtained reaction mixture (solution) was neutralized by adding 16% sulfuric acid, after which methyl isobutyl ketone was added to dissolve the reaction product. Thereafter, the water layer was separated and the obtained oil layer was rinsed in water. The oil layer was then distilled to obtain a concentrate. The concentrate was recrystallized by adding toluene and methanol, then filtered and dried to obtain 125 g of the target 4,4'-(1-trifluoromethyl-2-trifluoroethylidene) bis (3,5-dihydroxymethyl phenol) in refined state (94.1% purity as measured by high-speed liquid chromatography) as a white solid.

Melting point: 155.2°C

Mass spectrometry (LC-MS/AP): 456

Proton nuclear magnetic resonance spectrum (400 MHz, solvent: DMSO):

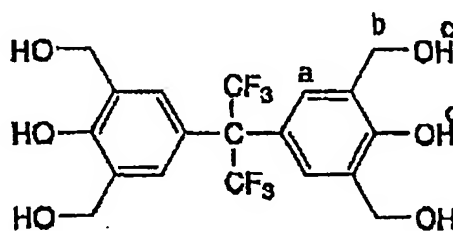
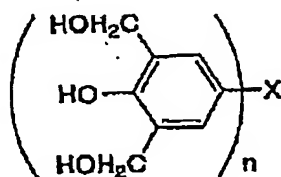


Table 4

δ (ppm)	Signal	Number of protons	Assignment
4.54	s	8	b
5.36	s	4	c
7.20	s	4	a
8.91	s	2	d

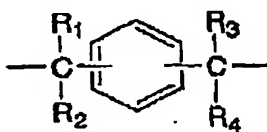
Claims

1. Hydroxymethyl-substituted polyfunctional phenol, expressed by the following structure:
General structure (I)



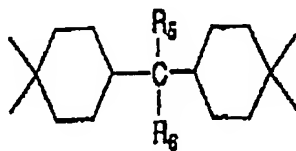
(I)

(wherein X represents: bivalent group (a) expressed by the following structure:
General structure (II)



(II)

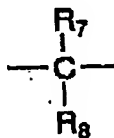
(wherein R_1 , R_2 , R_3 and R_4 each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4); quadrivalent group (b) expressed by the following structure:
General structure (III)



(III)

(wherein R_5 and R_6 each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);

or bivalent group (c) expressed by the following structure:
General structure (IV)

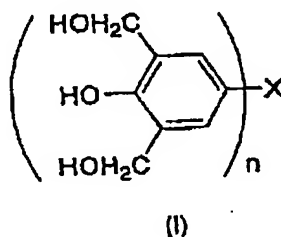


(IV)

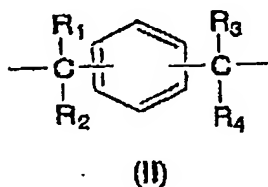
(wherein R_7 and R_8 each independently represent a hydrogen atom or a monofluoromethyl, difluoromethyl or trifluoromethyl group; however, R_7 and R_8 cannot be both hydrogen atoms); wherein n takes 2 when X is bivalent

group (a), takes 4 when X is quadrivalent group (b), or takes 2 when X is bivalent group (c)).

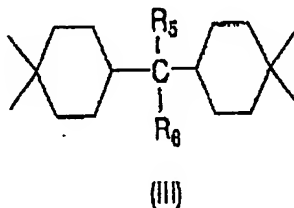
2. The hydroxymethyl-substituted polyfunctional phenol according to claim 1, wherein said R_1 , R_2 , R_3 and R_4 in General structure (II) each independently represent a methyl group, an ethyl group, a propyl group or a butyl group.
3. The hydroxymethyl-substituted polyfunctional phenol according to claim 1 or 2, wherein said R_5 and R_6 in General structure (III) each independently represent a methyl group, an ethyl group, a propyl group or a butyl group.
4. A method for producing a hydroxymethyl-substituted polyfunctional phenol, expressed by the following structure: General structure (I)



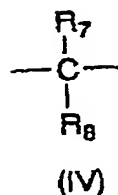
(wherein X represents: bivalent group (a) expressed by the following structure: General structure (II)



(wherein R_1 , R_2 , R_3 and R_4 each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);
quadrivalent group (b) expressed by the following structure:
General structure (III)

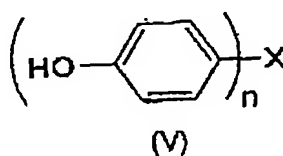


(wherein R_5 and R_6 each independently represent a hydrogen atom or an alkyl group with a carbon atom number of 1 through 4);
or bivalent group (c) expressed by the following structure:
General structure (IV)



(wherein R_7 and R_8 each independently represent a hydrogen atom or a monofluoromethyl, difluoromethyl or trifluoromethyl group; however, R_7 and R_8 cannot be both hydrogen atoms); wherein n takes 2 when X is bivalent group (a), takes 4 when X is quadrivalent group (b), or takes 2 when X is bivalent group (c)), said method comprising the steps of:

reacting multivalent phenol compounds expressed by general structure (V) below



(wherein X and n are the same as defined earlier) with 1 to 10 mols of formaldehyde per 1 mol of hydroxymethyl group contained in 1 mol of the target substance, in water or a mixed solvent consisting of water and organic solvent, in the presence of basic catalyst; and neutralizing the obtained reaction product to produce hydroxymethyl-substituted polyfunctional phenols.

5. Use of a hydroxymethyl-substituted polyfunctional phenol according to any of claims 1 to 3 for modifying phenol resins, as photo-resist material or for producing a polyphenol compound.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 03 00 7910

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	CHEMICAL ABSTRACTS, vol. 49, no. 10, 25 May 1955 (1955-05-25) Columbus, Ohio, US; S. SEITO ET AL: "Condensation reactions of formaldehyde with phenols. VI. Isomeric tetramethyloldihydroxydiphenylmethanes" column 6889; XP002244916 * abstract * & J. CHEM. SOC. JAPAN, IND. CHEM. SECT., vol. 56, 1953, pages 890-892, -----	1,4	C07C39/12 C07C39/367 C07C37/20
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			C07C
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 19 June 2003	Examiner Van Geyt, J
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

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